

ATTACHMENT 4-1

FROM :

FAX NO. : 6516338083

Aug. 24 1999 01:48PM FZ

DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE FOOD AND DRUG ADMINISTRATION		APPLICATION FOR A VARIANCE FROM 21 CFR 1040.11(c) FOR A LASER LIGHT SHOW, DISPLAY, OR DEVICE	FORM APPROVED, OMB NO. 0910-0075. EXPIRATION DATE: JUNE 28, 1998. FOR USE AT THE DATES INDICATED.														
			DOCKET NUMBER														
NOTE: No laser light show, projection system, or device may vary from compliance with 21 CFR 1040.11(c) in design or use without the approval of this application in accordance with 21 CFR 1010.4.																	
1. Check off applicable boxes and type or print the requested information. 2. Submit an original and four (4) copies. 3. NAME OF COMPANY <i>P.F. Entertainment, L.L.C.</i> 4. ADDRESS OF COMPANY (Include city code) (If P.O. box is used, include actual street address also) <i>100 Music Road, Pigeon Forge, TN 37863</i> 5. NAME AND TITLE OF RESPONSIBLE PERSON <i>Roger Reeder, Production Mgr.</i> 6. TELEPHONE NO. (Include area code) <i>423-428-7469</i> 7. DATE OF SUBMISSION <i>8-24-99</i>		INSTRUCTIONS 3. Mail your application to the Docket Management Branch (MDA-305), Food and Drug Administration, Room 6-52, 5600 Fishers Lane, Rockville, MD 20857 4. Enter docket number if assigned.															
8. THE APPLICANT REQUESTS THE VARIANCE TO BE IN EFFECT FOR A PERIOD OF _____ YEARS FROM THE DATE OF ISSUE. (In general, the Agency will approve a variance for only two years. If a longer period is requested, justification must be attached as part of the application.)																	
PRODUCT DESCRIPTION AND USE 9. LIST NAME AND/OR MODEL NUMBER(S) FOR THE LASER LIGHT SHOW(S) AND PROJECTOR(S) <i>Projector: Model CS Series -CS-12 / Name of Show: "Gita with Love"</i> 10. PRODUCT FOR WHICH A VARIANCE IS REQUESTED <input checked="" type="checkbox"/> Laser display device <input checked="" type="checkbox"/> Projector for a laser light show <input type="checkbox"/> A laser light show <input type="checkbox"/> Other (Specify) _____ 11. PROJECTORS ARE INTENDED FOR SALE, LEASE, OR LOAN TO OTHER LASER LIGHT SHOW PRODUCERS <input type="checkbox"/> 12. PRODUCT IS INTENDED FOR USE IN A <input type="checkbox"/> Planetarium or other dome projection structure <input type="checkbox"/> Theater <input type="checkbox"/> Motorized ballroom or meeting room <input type="checkbox"/> Store displays <input type="checkbox"/> Trade show or convention <input type="checkbox"/> Discotheque or night club <input type="checkbox"/> Pavilion <input type="checkbox"/> Indoor arena <input type="checkbox"/> Outdoor arena <input type="checkbox"/> Museum <input type="checkbox"/> Outdoor unenclosed area <input type="checkbox"/> Other (Specify) _____ 13. PRODUCT IS INTENDED TO BE USED <input checked="" type="checkbox"/> At only one (fixed) location <input type="checkbox"/> At a variety of (tour) locations <input type="checkbox"/> Other (Specify) _____				14. PRODUCT IS INTENDED TO BE USED AT ANY ONE LOCATION <input checked="" type="checkbox"/> More than 15 days <input type="checkbox"/> More than 5 but not more than 15 days <input type="checkbox"/> Less than 5 days 15. TOUR IS INTENDED TO RUN FOR <input type="checkbox"/> More than 6 months <input type="checkbox"/> 1-6 months <input type="checkbox"/> Less than one month <input type="checkbox"/> Not applicable (Not a tour) <input type="checkbox"/> Other (Specify) <i>Permanent Installation</i>													
		16. PRODUCT UTILIZES THE FOLLOWING LASER EFFECTS <input type="checkbox"/> Front screen projections <input type="checkbox"/> Rear screen projections <input type="checkbox"/> Holographic displays <input type="checkbox"/> Multiple reflection/refraction effects <input type="checkbox"/> Audience scanning (Also include scanning any accessible uncontrolled areas) <input checked="" type="checkbox"/> Reflections from stationary mirrors or mirrored surfaces (Beam Mirrors) <input type="checkbox"/> Stationary irradiation of rotating mirror balls, etc. <input type="checkbox"/> Scanning irradiation of rotating mirror balls, etc. <input type="checkbox"/> Fiber optic projections <input type="checkbox"/> Fog, smoke, or other scattering enhancement effects <input type="checkbox"/> Other (Specify) <i>Aerospace Aerial Beam effects</i>															
17. LASER RADIATION LEVELS <table border="1"> <thead> <tr> <th>LASER MEDIUM (e.g. Nd:YAG)</th> <th>WAVELENGTHS (nm)</th> <th>PEAK POWER (mW)</th> </tr> </thead> <tbody> <tr> <td><i>Nd: Yag</i></td> <td><i>532nm</i></td> <td><i>3 Watts</i></td> </tr> <tr> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td></td> <td></td> </tr> </tbody> </table>				LASER MEDIUM (e.g. Nd:YAG)	WAVELENGTHS (nm)	PEAK POWER (mW)	<i>Nd: Yag</i>	<i>532nm</i>	<i>3 Watts</i>								
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18. IF ANY LASER RADIATION IS PULSED OR SCANNED, GIVE THE PULSE DURATION AND RATE AND SCANNING FREQUENCY <i>N/A</i>																	
19. REASON FOR REQUESTING VARIANCE <input checked="" type="checkbox"/> Compliance with the limits of 21 CFR 1040.11(c) would restrict the intended use of the product because compliance would limit the output power to the extent that the desired effects would not be sufficiently visible <input type="checkbox"/> Other or additional explanation (Specify) _____																	

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VAR 1

FROM :

FAX NO. : 6516338883

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11. MANNER IN WHICH IT IS PROPOSED TO DEVIATE FROM THE REQUIREMENTS OF THE APPLICABLE STANDARD

- It is proposed to deviate from the provisions of 21 CFR 1040.11(c) in that the accessible emission levels would exceed the accessible emission limits specified in 21 CFR 1040.11(c).
- It is proposed to deviate from the provision of 21 CFR 1040.11(c) as follows:

12. ADVANTAGES TO BE DERIVED FROM SUCH DEVIATION

- Laser light shows and displays are popular media in entertainment and the arts. Use of power levels in excess of the limits imposed by 21 CFR 1040.11(c) is necessary to achieve the required effects in these media.
- Other or additional advantages (describe and explain)

13. EXPLAIN THE ALTERNATE MEANS OF RADIATION PROTECTION TO BE PROVIDED. (Check as many boxes as apply. In Item 7d "Remarks," justify any boxes not checked, using additional sheets as necessary. State any other means of radiation protection that will be used.)

- a. All laser projectors, systems, shows, and projectors will be certified to comply with 21 CFR 1040.10 and the conditions of this variance and will be reported as required by 21 CFR 1042.10 and 1042.12 using the reporting guides provided for such purpose. These actions will be accomplished prior to any introduction into commerce.
- b. Effects not specifically indicated in this variance application will not be performed. No other effects will be added until an amendment to the variance has been obtained and the required reports or supplements, as applicable, have been submitted.
- c. Scanning, projection, or reflection of laser and collateral radiation (light show effects) into audience or other accessible uncontrolled areas will not be permitted except for diffuse reflections produced by the atmosphere, added atmospheric scattering media, and target effects.
- d. Laser radiation levels, in excess of the limits of Class I will not be permitted at any point less than 3.0 meters above any surface upon which persons other than operators, performers, or employees are permitted to stand or 2.5 meters below or in lateral separation from any place where such persons are permitted to be. Operators, performers, and employees will not be required or allowed to view radiation above the limits of Class I or be exposed to radiation above the limits specified in 21 CFR 1040.11(c).
- e. Any product which relies on scanning to scan access, exposure, or product class limits will incorporate a scanning safeguard system which directly senses scanner motion and which will react fast enough to prevent exceeding the applicable limit.
- f. All laser light shows shall be under the direct and personal control of trained, competent operator(s). The operator(s) will:
- (1) immediately terminate the emission of light show radiation in the event of any unsafe condition;
 - (2) Be located where all beam paths can be directly observed at all times; and
 - (3) Be an employee of the variance holder, who will be responsible for the training and the conduct of the operator.
- g. The maximum laser projector output power will not exceed the level required to obtain the intended effects.
- h. The projection system (i.e., projector and all other components used to produce the lighting effects) will be securely mounted or immobilized to prevent accidental movement or misalignment. Beam masking will be provided as an inherent part of the system design to prevent overfilling of screens, beam stage, targets, etc.
- i. Laser projectors will not be delivered to any other party under an agreement of sale, lease, or loan unless and until the recipient demonstrates that they have a variance in effect at the time of delivery that permits them to produce laser light shows incorporating such projector(s).
- j. In addition to the requirements of 21 CFR 1040.10(n), the manufacturer of laser projector systems will provide to parties who purchase, lease, or borrow the equipment, adequate user instructions for safe installation and operation which explain the responsibility of the recipient as an independent light show manufacturer to submit the required reports and apply for and obtain a variance from FDA prior to introduction into commerce of any laser light show.
- k. The requirements of 21 CFR 1042.30(a)(1) and (2) will be accomplished through the use of written procedures for setup, alignment, testing, and performance of each show. These procedures will be in sufficient detail to ensure compliance with 21 CFR 1040.10, the conditions of this variance, and the control of access to radiation areas using the procedures described in the ANSI Z136.1 standard for the safe use of lasers (American National Standards Institute, 747 Madison Avenue, New York, NY 10017) or any other equivalent user consensus standard and, where applicable, state or local requirements. Laser radiation areas which can contain radiation levels above the limits specified in 21 CFR 1040.11(c) will be clearly identified by the posting of warning signs and/or restricting access through physical means (such as pressure switches, photo cells, barriers, guards, etc.). These requirements apply to temporary areas (such as during set up and alignment procedures) and to final or permanent areas. The variance holder will retain the records of these procedures and the results of all tests as required by 21 CFR 1042.31. A copy of the variance application, the approval letter, current procedures, and records relating to each particular show will be with the operator or other responsible individual and will be made available for inspection by FDA and other responsible authorities.

FROM :

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Aug. 24 1999 02:52PM PT

- Advance written notification will be made as early as possible to appropriate federal, state, and local authorities (including show industry) with dates and locations clearly and completely identified, and a brief description of the proposed effects including a statement of the maximum power output intended. Such notifications will be made, but not necessarily be limited, to:
- (1) The Center for Devices and Radiological Health, Office of Compliance (HFD-212), 2058 Gaither Road, Rockville, MD 20850, providing the initial and closing notice for fixed installations and the itinerary for mobile shows. In addition, unless all aspects of each show have been reported and accession numbers clearly referenced, each notice will include detailed descriptions of each show and a listing of all effects to be performed in sufficient detail to confirm compliance with the regulations and this variance.
 - (2) The Federal Aviation Administration (FAA) for any projections into open airspace of any time (i.e., including set up, alignment, repositioning, performance, etc.). If the FAA objects to any laser effect, the objections will be resolved and any conditions requested by FAA will be adhered to. If these conditions cannot be met, the objectionable effects will be deleted from the show.
 - (3) State and local aviation control units/agencies for all shows to be performed within their jurisdictions. All requirements of state and local law will be satisfied and any objection raised by local authorities will be resolved or the effects deleted. (A list of federal and state offices is available from the Center for Devices and Radiological Health upon request.)

14. REMARKS

CERTIFICATION

I CERTIFY that all of the above information and statements are true, complete, and correct to the best of my knowledge and acknowledge that any variance application may be denied or my variance may be revoked if any application is found to be false, misleading, or incorrect in any material way. I have submitted and will submit all reports required by 21 CFR 1002.10 and 1002.12 on the laser equipment and show(s). I further understand that I may be required by regulation or by the Director, Center for Devices and Radiological Health to supply such other information as may be necessary to evaluate and act on this application.

15. SIGNATURE

GSA FPMR 3147 (b)(5)

16. NAME (Type or Print)

Roger M. Redden

17. TITLE

TECHNICAL DIRECTOR
LASER SAFETY OFFICER

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